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Dated

24 September 2004

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1/77

9 OCT 2003

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The Patent Office

Cardiff Road
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South Wales
NP10 8QQ

1. Your Reference

PB60535P

2. Patent application number

(The Patent office will fill in this part)

0323701.3

100CT03 E843510-1 D02029

PD1/7700 0.00-0323701.3

3. Full name, address and postcode of the or of each applicant (underline all surnames)

GLAXO GROUP LIMITED
GLAXO WELLCOME HOUSE
BERKELEY AVENUE
GREENFORD
MIDDLESEX
UB6 ONN
GB

Patents ADP number (if you know it)

If the applicant is a corporate body, give the country/state of its corporation

00473587008
GB

4 Title of the invention

FORMULATIONS

5 Name of your agent (if you have one)

"Address for service" in the United Kingdom to which all correspondence should be sent (including the postcode)

GLAXOSMITHKLINE
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BRENTFORD, MIDDLESEX
TW8 9GS, GB

Patents ADP number (if you know it)

08072555006

6. Priority: Complete this section if you are declaring priority from one or more earlier patent applications, filed in the last 12 months

Country

Priority application number
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Date of Filing
(day / month / year)

7. Divisionals: etc Complete this section only if this application is a divisional application or resulted from an entitlement dispute (see note f)

Number of earlier UK application

Date of filing
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8. Is a Patents Form 7/77 (Statement of inventorship and of right to grant of a patent) required in support of this request?

YES

Answer YES if:

a) any applicant named in part 3 is not an inventor, or

b) there is an inventor who is not named as an applicant, or

c) any named applicant is a corporate body

Otherwise answer NO See note (d)

Patents Form 1/77

9. Accompanying documents: A patent application must include a description of the invention. Not counting duplicates, please enter the number of pages of each item accompanying this form:

Continuation sheet of this form

Description	14
Claim(s)	-
Abstract	-
Drawing(s)	-

10. If you are also filing any of the following, state how many against each item

Priority Documents

Translations of priority documents

Statement of inventorship and right to grant of a patent (*Patents Form 7/77*)

Request for preliminary examination and search (*Patent Form 9/77*)

Request for substantive examination (*Patent Form 10/77*)

Any other documents (*please specify*)

11. I/We request the grant of a patent on the basis of this application

Signature(s)

Judith Pritchard

**JUDITH PRITCHARD
AGENT FOR THE APPLICANTS**

Date: 9 October 2003

12. Name and daytime telephone number and e-mail address, if any, of person to contact in the United Kingdom
- KATHERINE EVANS 01438 768611**

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Formulations

The present invention relates to novel pharmaceutical formulations of particulate drug such as beta-agonists and/or anti-inflammatory steroids in hydrofluoroalkane propellants with the carboxylic acid surfactant compounds particularly [(7,7,8,8,8-pentafluorooctyl)oxy]acetic acid, preparation of said formulation and their use in therapy, and the use of said surfactant in reducing the variability in the content uniformity of formulations or in providing enhanced FPM in 1,1,1,2-tetrafluoroethane (134a) and/or 1,1,1,2,3,3,3-heptafluoro-n-propane (227) suspension formulations.

The use of aerosols to administer medicaments has been known for several decades. Such aerosols generally comprise the medicament, one or more chlorofluorocarbon propellants and either a surfactant or a co-solvent, such as ethanol. Historically the most commonly used aerosol propellants for medicaments have been propellant 11 (CCl_3F) and/or propellant 114 ($\text{CF}_2\text{ClCF}_2\text{Cl}$) with propellant 12 (CCl_2F_2). However these propellants are now believed to provoke the degradation of stratospheric ozone and there is thus a need to provide aerosol formulations for medicaments which employ so called "ozone-friendly" propellants.

A class of propellants which are believed to have minimal ozone-depleting effects in comparison to conventional chlorofluorocarbons comprise fluorocarbons and hydrogen-containing chlorofluorocarbons, and a number of medicinal aerosol formulations using such propellant systems are disclosed in, for example, EP0372777, WO91/04011, WO91/11173, WO91/11495 and WO91/14422. These applications are all concerned with the preparation of pressurised aerosols for the administration of medicaments and seek to overcome the problems associated with the use of the new class of propellants, in particular the problems of stability associated with the pharmaceutical formulations prepared. The applications all propose the addition of one or more of adjuvants such as alcohols, alkanes, dimethyl ether, surfactants (including fluorinated and non-fluorinated surfactants, carboxylic acids, polyethoxylates etc) and even conventional chlorofluorocarbon propellants in small amounts intended to minimise potential ozone damage.

WO92/00061 discloses non-fluorinated surfactants for use with fluorocarbon propellants.

It is essential that the prescribed dose of aerosol medication delivered from the MDI to the patient consistently meets the specifications claimed by the manufacturer and comply with the requirements of the FDA and other regulatory authorities. That is, every dose dispensed from the can must be the same within close tolerances. Therefore it is important that the formulation be substantially homogenous throughout

the canister and the administered dose at the time of actuation of the metering valve and remains substantially the same even after storage. Thus the uniformity of the dose dispensed through the life of the device is vitally important.

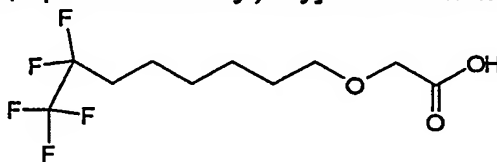
In the case of suspension formulations, to control aggregation of fine particles, and thereby influence the dispersability of the suspension it is well established in the art that fluorinated surfactants may be used to stabilise micronised drug suspensions in fluorocarbon propellants such as 134a or 227, see for example US4352789, US5126123, US5376359, US application 09/580008, WO91/11173, WO91/14422, WO92/00062 and WO96/09816.

The problem of aggregation of the particulate drug may be manifest as a drop in fine particle mass (FPM) after storage. The FPM is a measure of the dose dispensed which has the potential to reach the therapeutic portion of the lung. Thus a drop in FPM means the therapeutically effective amount of drug available to the patient is reduced which is undesirable and may ultimately be dangerous. This problem is particularly acute when the dose due to be dispensed is low, which is the case for certain potent drugs such as long acting beta agonists.

Furthermore, it is desirable to have a mass median aerodynamic diameter (MMAD) of particles, is within a controlled predetermined range to maximise the therapeutic effect of the dose dispensed. The MMAD has proved difficult to control, in many instances for formulations of the new propellants.

Suspension formulations which are not adequately stabilised result in high levels of drug deposition, for example, on the canister walls or on components of the metered dose inhaler, such as the valve components including the metering chamber, seals or the like. This deposition may not only result in drug loss thereby reducing the total drug content of the canister available to patient but can also adversely affect the functioning of the device, resulting in the valve sticking, orifices becoming blocked, caking of drug which may work free at a latter point and increase the dose given to the patient in an unpredictable way. Furthermore, expensive modifications to the canister and/or valve may be required to deal with this deposition.

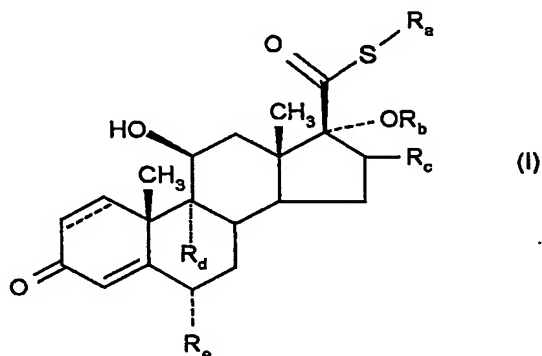
While the use of surfactants, including certain polyoxyethylene surfactants has been suggested, for example in WO95/15151, US 5676931 and WO92/00061, the inventors have now found [(7,7,8,8,8-pentafluorooctyl)oxy]acetic acid which has the formula



has particularly good surfactant properties in 1,1,1,2-tetrafluoroethane and/or 1,1,1,2,3,3,3-heptafluoro-n-propane in comparison to the general class of surfactants, and is particularly useful when employed in pharmaceutical aerosol formulations for administering a particulate medicament of formula (I) and/or of formula (II), the structures of which are given below, to the lungs thereby ameliorating or solving one or more of the above problems.

Thus the invention provides pharmaceutical aerosol formulations for administering a particulate medicament to the lungs comprising:

i) a therapeutic effective amount of particulate drug selected from a compound of formula (I)



wherein

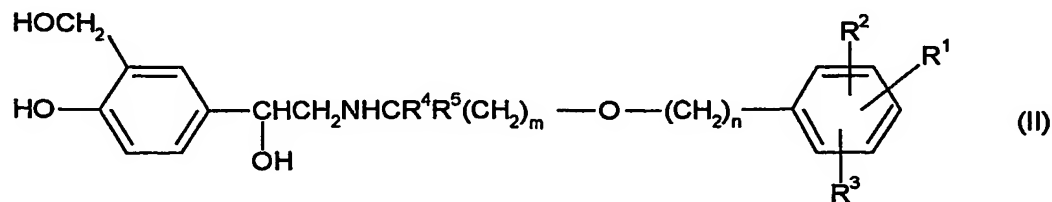
R_a represents C_{1-6} alkyl or C_{1-6} haloalkyl;

R_b represents $-C(=O)-$ aryl or $-C(=O)-$ heteroaryl;

R_c represents hydrogen, methyl (which may be in either the α or β configuration) or methylene;

R_d and R_e are the same or different and each represents hydrogen or halogen; and

----- represents a single or a double bond and salts and solvates thereof, for example, as disclosed WO02/12265 and WO02/12266, and/or a compound of formula (II)



or a salt, solvate, or physiologically functional derivative thereof, wherein:

m is an integer of from 2 to 8;

n is an integer of from 3 to 11;

with the proviso that m + n is 5 to 19;

R¹ is -XSO₂NR⁶R⁷

wherein X is -(CH₂)_p- or C₂₋₆ alkenylene;

R⁶ and R⁷ are independently selected from hydrogen, C₁₋₆alkyl,

C₃₋₇cycloalkyl, C(O)NR⁸R⁹, phenyl, and phenyl (C₁₋₄alkyl)-,

or R⁶ and R⁷, together with the nitrogen to which they are bonded, form a 5-, 6-, or 7-membered nitrogen containing ring,

and R⁶ and R⁷ are each optionally substituted by one or two groups selected from halo,

C₁₋₆alkyl, C₁₋₆haloalkyl, C₁₋₆alkoxy, hydroxy-substituted C₁₋₆alkoxy, -CO₂R⁸, -SO₂NR⁸R⁹,

-CONR⁸R⁹, -NR⁸C(O)R⁹, or a 5-, 6- or 7-membered heterocyclic ring;

R⁸ and R⁹ are independently selected from hydrogen, C₁₋₆alkyl,

C₃₋₆cycloalkyl, phenyl, and phenyl (C₁₋₄alkyl)-; and

p is an integer of from 0 to 6;

R² and R³ are independently selected from hydrogen, C₁₋₆alkyl, C₁₋₆alkoxy, halo, phenyl, and C₁₋₆haloalkyl; and

R⁴ and R⁵ are independently selected from hydrogen and C₁₋₄alkyl with the proviso that the total number of carbon atoms in R⁴ and R⁵ is not more than 4, as disclosed in WO02/066422;

(ii) a propellant selected from the group comprising 1,1,1,2-tetrafluoroethane or 1,1,1,2,3,3,3-heterofluoro-n-propane and mixtures thereof; and

(iii) a surfactant [(7,7,8,8,8-pentafluorooctyl)oxy]acetic acid.

Advantageously, the surfactant compound of the present invention have good surfactant properties, such as reducing the mean length weighted diameter of suspension formulations, stabilising FPM and/or giving good content uniformity performance, of formulations of the above medicaments in 134a and/or 227 whilst avoiding toxic effects observed for certain perfluorinated surfactant compounds. Usually the best surfactants in 134a and 227 are perfluorinated compounds with a large number of perfluorinated carbon atoms, which unfortunately have a tendency to bioaccumulate. Thus there is an inherent conflict between good surfactant properties and minimising toxic effects. The present invention is advantageous in terms of improving the stability of the aerosol formulation by reducing drug deposition, increasing shelf life and the like.

Whilst not wishing to be bound by theory it is thought that the properties of the surfactant are particularly well matched to those of the medicaments employed in the

formulations of the present invention, for example, in respect of their pKa and thereby provide good stabilising effects.

In a first aspect the invention provides a pharmaceutical aerosol formulation wherein the medicament comprises 3-(4-[[6-((2*R*)-2-hydroxy-2-[4-hydroxy-3-(hydroxymethyl)phenyl]ethyl) amino)hexyl]oxy}butyl) benzenesulfonamide or 3-(3-[[7-((2*R*)-2-hydroxy-2-[4-hydroxy-3-(hydroxymethyl)phenyl]ethyl)amino]heptyl]oxy}propyl) benzenesulfonamide.

In a second aspect the invention provides a pharmaceutical aerosol formulation wherein the medicament comprises 6 α , 9 α -difluoro-17 α -[(2-furanylcarbonyl)oxy]-11 β -hydroxy-16 α -methyl-3-oxo-androsta-1,4-diene-17 β -carbothioic acid S-fluoromethyl ester or 6 α ,9 α -difluoro-11 β -hydroxy-16 α -methyl-17 α -[(4-methyl-1,3-thiazole-5-carbonyl)oxy]-3-oxo-androsta-1,4-diene-17 β -carbothioic acid S-fluoromethyl ester.

In a third aspect the present invention provides a pharmaceutical aerosol formulation wherein the medicament comprises 3-(4-[[6-((2*R*)-2-hydroxy-2-[4-hydroxy-3-(hydroxymethyl)phenyl]ethyl) amino)hexyl]oxy}butyl) benzenesulfonamide and 6 α , 9 α -difluoro-17 α -[(2-furanylcarbonyl)oxy]-11 β -hydroxy-16 α -methyl-3-oxo-androsta-1,4-diene-17 β -carbothioic acid S-fluoromethyl ester or 6 α ,9 α -difluoro-11 β -hydroxy-16 α -methyl-17 α -[(4-methyl-1,3-thiazole-5-carbonyl)oxy]-3-oxo-androsta-1,4-diene-17 β -carbothioic acid S-fluoromethyl ester.

In a fourth aspect the present invention provides a pharmaceutical aerosol formulation wherein the medicament comprises 3-(3-[[7-((2*R*)-2-hydroxy-2-[4-hydroxy-3-(hydroxymethyl)phenyl]ethyl)amino]heptyl]oxy}propyl) benzenesulfonamide and 6 α , 9 α -difluoro-17 α -[(2-furanylcarbonyl)oxy]-11 β -hydroxy-16 α -methyl-3-oxo-androsta-1,4-diene-17 β -carbothioic acid S-fluoromethyl ester or 6 α ,9 α -difluoro-11 β -hydroxy-16 α -methyl-17 α -[(4-methyl-1,3-thiazole-5-carbonyl)oxy]-3-oxo-androsta-1,4-diene-17 β -carbothioic acid S-fluoromethyl ester.

In this specification when a medicament is referred to also are salts and solvates thereof such as pharmacologically acceptable salts.

Where the medicament employed is 3-(4-[[6-((2*R*)-2-hydroxy-2-[4-hydroxy-3-(hydroxymethyl)phenyl]ethyl) amino)hexyl]oxy}butyl) benzenesulfonamide preferably it is employed as the cinnamate salt.

In an alternative aspect the invention extends to formulations of comprising [(7,7,8,8,8-pentafluorooctyl)oxy]acetic acid, a propellant such as 134a and/or 227 and a particulate drug selected from the group comprising:

3-(3-[[7-((2R)-2-hydroxy-2-[4-hydroxy-3-hydroxymethyl]phenyl]ethyl)-amino]heptyl]oxy}propyl)benzenesulfonamide,
4-((1R)-2-[(6-{2-[(2,6-dichlorobenzyl)oxy]ethoxy}hexyl)amino]-1-hydroxyethyl)-2-(hydroxymethyl)phenol,
N-(3,5-dichloropyridin-4-yl)-2-[1-(4-fluorobenzyl)-5-hydroxyindol-3-yl]-2-oxoacetamide,
a compound of formula (II) or claim as disclosed in WO01/42193,
a compound of formula (I) as disclosed in WO03/042160,
a compound of formula (I) as disclosed in WO03/042164 and combinations thereof.
and/or

[(7,7,8,8,8-Pentafluorooctyl)oxy]acetic acid can be prepared as described in WO03/013610.

Use of said surfactant compounds for the preparation of formulations according to the present invention results in effective suspension stabilisation at low concentrations relative to the amount of medicament. Thus, the amount of the surfactant employed is desirably in the range of 0.05% to 20% w/w, particularly 0.5% to 10% w/w, more particularly 0.5% to 5% w/w, relative to the medicament.

The particle size of the particulate (e.g. micronised) medicament should be such as to permit inhalation of substantially all of the medicament into the lungs or nasal cavity upon administration of the aerosol formulation and will thus be less than 100 microns, desirably less than 20 microns, and preferably will have a MMAD in the range 1-10 microns, e.g. 1-5 microns.

The final aerosol formulation desirably contains 0.005-10% w/w, preferably 0.005 - 5% w/w, especially 0.01-1.0% w/w, of medicament relative to the total weight of the formulation.

Preferably a single propellant is employed, for example, 1,1,1,2-tetrafluoroethane or 1,1,1,2,3,3,3-heptafluoro-n-propane, especially 1,1,1,2-tetrafluoroethane.

It is desirable that the formulations of the invention contain no components which may provoke the degradation of stratospheric ozone. In particular it is desirable that the formulations are substantially free of chlorofluorocarbons such as CCl₃F, CCl₂F₂ and CF₃CCl₃.

If desired the propellant may additionally contain a volatile adjuvant such as a saturated hydrocarbon, for example, propane, n-butane, isobutane, pentane and isopentane or a dialkyl ether, for example, dimethyl ether. In general, up to 50% w/w of the propellant may comprise a volatile hydrocarbon, for example 1 to 30% w/w.

However, formulations which are substantially free of volatile adjuvants are preferred. In certain cases, it may be desirable to include appropriate amounts of water, which can be advantageous in modifying the dielectric properties of the propellant.

Polar adjuvants which may if desired, be incorporated into the formulations according to the present invention include, for example, C₂₋₆aliphatic alcohols and polyols such as ethanol, isopropanol and propylene glycol and mixtures thereof. Preferably ethanol will be employed. In general only small quantities (e.g. 0.05 to 3.0% w/w) of polar adjuvants are required and the use of quantities in excess of 5% w/w may disadvantageously tend to dissolve the medicament. Formulations preferably contain less than 1% w/w, for example, about 0.1% w/w of polar adjuvant. Most preferably the formulations according to the invention are substantially free of polar adjuvant. Polarity may be determined, for example, by the method described in European Patent Application Publication No. 0327777.

In addition to the surfactant, the formulations according to the present invention may optionally contain one or more further ingredients conventionally used in the art of pharmaceutical aerosol formulation. Such optional ingredients include, but are not limited to, taste masking agents, sugars, buffers, antioxidants, water and chemical stabilisers.

A particularly preferred embodiment of the invention provides a pharmaceutical aerosol formulation consisting essentially of one or more particulate medicament(s) of formula (I) and/or formula (II), or other medicament disclosed in this specification or combination thereof, one or more of said propellants and [(7,7,8,8,8-pentafluorooctyl)oxy]acetic acid.

The invention also extends to formulations as described already which consist rather than comprise said elements.

A further embodiment of the invention is a sealed container capable of withstanding the pressure required to maintain the propellant as a liquid, such as a metered dose inhaler, containing therein one of the aerosol formulation as described above.

The term "metered dose inhaler" or MDI means a unit comprising a can, a secured cap covering the can and a formulation metering valve situated in the cap. MDI system includes a suitable channelling device. Suitable channelling devices comprise for example, a valve actuator and a cylindrical or cone-like passage through which medicament may be delivered from the filled canister via the metering valve to the nose or mouth of a patient such as a mouthpiece actuator.

MDI canisters generally comprise a container capable of withstanding the vapour pressure of the propellant used such as a plastic or plastic-coated glass bottle or preferably a metal can, for example, aluminium or an alloy thereof which may optionally be anodised, lacquer-coated and/or plastic-coated (e.g. incorporated herein by reference WO96/32099 wherein part or all of the internal surfaces are coated with one or more fluorocarbon polymers optionally in combination with one or more non-fluorocarbon polymers), which container is closed with a metering valve. The cap may be secured onto the can via ultrasonic welding, screw fitting or crimping. MDIs taught herein may be prepared by methods of the art (e.g., see Byron, above and WO/96/32099). Preferably the canister is fitted with a cap assembly, wherein a drug metering valve is situated in the cap, and said cap is crimped in place.

Advantageously formulations according to the present invention obviate the need for the additional processing of the canisters and/or component by, for example, coating which ultimately leads to cost saving, especially when manufacturing in bulk.

The metering valves are designed to deliver a metered amount of the formulation per actuation and incorporate a gasket to prevent leakage of propellant through the valve. The gasket may comprise any suitable elastomeric material such as, for example, low density polyethylene, chlorobutyl, black and white butadiene-acrylonitrile rubbers, butyl rubber and neoprene. Suitable valves are commercially available from manufacturers well known in the aerosol industry, for example, from Valois, France (e.g. DF10, DF30, DF60), Bepak plc, UK (e.g. BK300, BK357) and 3M-Neotechnic Ltd, UK (e.g. SpraymiserTM).

The formulations of the invention may be prepared by dispersal of a compound of formula (I) and/or (II) or other medicament as appropriate and the chosen surfactant compound in the selected propellant in an appropriate container, for example, with the aid of sonication or a high-shear mixer. The process is desirably carried out under controlled humidity conditions.

Alternatively the surfactant compound may be prespiked into an empty canister before the cap and valve are secured in place.

A further aspect of this invention comprises a process for filling the said formulation into MDIs.

Conventional bulk manufacturing methods and machinery well known to those skilled in the art of pharmaceutical aerosol manufacture may be employed for the preparation of large scale batches for the commercial production of filled canisters. Thus, for example, in one bulk manufacturing method a metering valve is crimped onto an

aluminium can to form an empty canister. The particulate medicament is added to a charge vessel and liquefied propellant is pressure filled through the charge vessel into a manufacturing vessel, together with liquefied propellant containing the surfactant. The drug suspension is mixed before recirculation to a filling machine and an aliquot of the drug suspension is then filled through the metering valve into the canister.

In an alternative process, an aliquot of the liquefied formulation is added to an open canister under conditions which are sufficiently cold to ensure formulation does not vaporise, and then a metering valve crimped onto the canister.

Typically, in batches prepared for pharmaceutical use, each filled canister is check-weighed, coded with a batch number and packed into a tray for storage before release testing.

Each filled canister is conveniently fitted into a suitable channelling device prior to use to form a metered dose inhaler system for administration of the medicament into the lungs or nasal cavity of a patient.

The chemical and physical stability and the pharmaceutical acceptability of the aerosol formulations according to the invention may be determined by techniques well known to those skilled in the art. Thus, for example, the chemical stability of the components may be determined by HPLC assay, for example, after prolonged storage of the product. Physical stability data may be gained from other conventional analytical techniques such as, for example, by leak testing, by valve delivery assay (average shot weights per actuation), by dose reproducibility assay (active ingredient per actuation) and spray distribution analysis.

The suspension stability of the aerosol formulations according to the invention may be measured by conventional techniques, for example, by measuring flocculation size distribution using a back light scattering instrument or by measuring particle size distribution by cascade impaction or by the "twin impinger" analytical process. As used herein reference to the "twin impinger" assay means "Determination of the deposition of the emitted dose in pressurised inhalations using apparatus A" as defined in British Pharmacopoeia 1988, pages A204-207, Appendix XVII C. Such techniques enable the "respirable fraction" of the aerosol formulations to be calculated. One method used to calculate the "respirable fraction" is by reference to "fine particle fraction" which is the amount of active ingredient collected in the lower impingement chamber per actuation expressed as a percentage of the total amount of active ingredient delivered per actuation using the twin impinger method described above.

Metered dose inhalers are designed to deliver a fixed unit dosage of medicament per actuation or "puff", for example, in the range of 10 to 5000 micrograms of medicament per puff.

Administration of medicament may be indicated for the treatment of mild, moderate, severe acute or chronic symptoms or for prophylactic treatment. It will be appreciated that the precise dose administered will depend on the age and condition of the patient, the particular particulate medicament used and the frequency of administration and will ultimately be at the discretion of the attendant physician. When combinations of medicaments are employed the dose of each component of the combination will in general be that employed for each component when used alone. Typically, administration may be one or more times, for example, from 1 to 8 times per day, giving for example 1, 2, 3 or 4 puffs each time.

An appropriate dosing regime for other medicaments will be known or readily available to persons skilled in the art.

The use of compounds [(7,7,8,8,8-pentafluorooctyl)oxy]acetic acid to enhance the FPM or reduce the variability in the content uniformity, for example, by reducing the relative standard deviation (RDS).

The following non-limiting examples serve to illustrate the invention.

Examples

Example 1

Standard 8 mL MDI cans, coated with a polymer blend of PTFE and polyether sulfone, were pre-spiked with 0.6 mg of [(7,7,8,8,8-pentafluorooctyl)oxy]acetic acid, the valves crimped in place, and a suspension of about 6 mg 3-(4-[[6-((2*R*)-2-hydroxy-2-(4-hydroxy-3-(hydroxymethyl) phenyl)ethyl] amino)hexyl]oxy}butyl) benzenesulfonamide (DRUG 1) in about 12 g P134a was filled through the valve.

Example 2

Standard 8 mL MDI cans, coated with a polymer blend of PTFE and polyether sulfone, were pre-spiked with 0.6 mg of [(7,7,8,8,8-pentafluorooctyl)oxy]acetic acid, the valves crimped in place, and a suspension of about 6 mg of 6 α , 9 α -difluoro-17 α -(2-furanylcarbonyl)oxy]-11 β -hydroxy-16 α -methyl-3-oxo-androsta-1,4-diene-17 β -carbothioic acid *S*-fluoromethyl ester (DRUG 2) in about 12 g of P134a was filled through the valve.

Example 3

Standard 8 mL MDI cans, coated with a polymer blend of PTFE and polyether sulfone, were pre-spiked with 0.7 mg of [(7,7,8,8,8-pentafluorooctyl)oxy]acetic acid, the valves crimped in place, and a suspension of about 3 mg of 3-(4-[[6-((2R)-2-hydroxy-2-[4-hydroxy-3-(hydroxymethyl) phenyl]ethyl] amino)hexyl]oxy}butyl) benzenesulfonamide (DRUG 1) and 4 mg of 6 α , 9 α -difluoro-17 α -[(2-furanylcarbonyl)oxy]-11 β -hydroxy-16 α -methyl-3-oxo-androsta-1,4-diene-17 β -carbothioic acid S-fluoromethyl ester (DRUG 2) in about 12 g of P134a was filled through the valve.

EXPERIMENTAL DATA**MEAN LENGTH WEIGHTED EQUIVALENT DIAMETER**

Mean length weighted equivalent diameter of drug suspensions in propellant P134a.

Mean length weighted equivalent diameter of suspensions was determined by image analysis (Galai CIS-100 image analyser) . It represents the diameter a circle of equivalent area to the object under analysis (weighted by particle diameter in the distribution)

Table 1 shows mean particle size data determined by image analysis using a Galai CIS-100 particle size analyser for sample formulations prepared as described above. In this measurement, particle size is represented as the equivalent diameter of a circle of equal area to the object. The mean is the average of 4 determinations. The particle size measurement was obtained by transferring the suspensions to a pressurised cell, and video-imaging the sample under shear via a microscope objective.

The equivalent diameter is defined as the diameter of a circle of equal area to the object.

$$\text{Equivalent Diameter} = \sqrt{\frac{\text{Area}}{\pi}}$$

The mean equivalent diameter can be weighted by number, length or volume.
e.g. For three particles with equivalent diameters of x, y and z:

$$\text{Mean Number weighted diameter} = \left(\frac{1}{3}\right)x + \left(\frac{1}{3}\right)y + \left(\frac{1}{3}\right)z$$

$$\text{Mean Length weighted diameter} = \left(\frac{x}{x+y+z}\right)x + \left(\frac{y}{x+y+z}\right)y + \left(\frac{z}{x+y+z}\right)z$$

The data shows that the surfactant in formulations according to the invention has suspension stabilising properties thereby discouraging flocculation of drug particles.

This is seen by the marked reduction in average particle size ("mean length") when the surfactant is incorporated into the formulation.

TABLE 1

Sample	Mean Length Weighted Equivalent Diameter (μm)
CONTROL (Surfactant-free)	$23.8 \pm 3.2 \mu\text{m}$
Example 1	$8.1 \pm 0.9 \mu\text{m}$
CONTROL (Surfactant-free)	$14.8 \pm 2.8 \mu\text{m}$
Example 2	$11.1 \pm 1.7 \mu\text{m}$

FPM

Table 2 shows data relating to the FPM (the sum of stages 3 to 5) obtained using an Anderson Cascade Impactor stack. Data were obtained at the beginning of use of the device. Controls were prepared corresponding to each of the samples but omitting the surfactant. The analysis of aerosol formulations using such stacks is well known to person skilled in the art. The data is shown as absolute FPM in μg and percentage FPM (in brackets) which expresses absolute FPM as a percentage of the total ex-valve emitted dose. This data is given for an initial timepoint and then after storage for 12 weeks at 40°C and 75% relative humidity.

Table 2 shows samples containing surfactant show an increase in the absolute value of the FPM fraction in most cases. This indicates that a greater proportion of the dose will be available to reach the therapeutic target of in the lung, which is desirable. Furthermore, it can be seen that the FPM in for Example 2 is stabilised by the presence of the surfactant.

TABLE 2

SAMPLE	FPM μg (FPM as % of dose emitted ex-valve)	
	Initial	12 Weeks 40°C/75% RH
CONTROL 1 (Surfactant-free)	6.9 μg (32.9 %)	4.9 μg (22.8%)
Example 1	11.2 μg (43.4 %)	8.8 μg (35.0 %)
CONTROL 2 (Surfactant-free)	7.9 μg (35.5 %)	6.7 μg (31.5%)
Example 2	7.2 μg (33.7 %)	7.2 μg (31.8%)
CONTROL DRUG 1 (Surfactant-free)	3.1 μg (31.0 %)	3.8 μg (38.7 %)
DRUG1	4.3 μg (36.5 %)	5.1 μg (46.1 %)
CONTROL DRUG 2 (Surfactant-free)	6.7 μg (30.0 %)	7.3 μg (30.1 %)
DRUG 2	8.5 μg (33.2 %)	9.0 μg (36.0 %)

CONTENT UNIFORMITY

The content uniformity of the formulation, the preparation of which is described above, was assessed by dose through use testing. Testing was performed on 10 cans/inhalers at "beginning of use" (BoU) and "end of use" (EoU), after storage for 12 weeks at 40°C and 75% relative humidity. After each inhaler had been primed (4 shots fired to waste), actuations 1 and 2 (BoU) were collected. The next 116 actuations of each inhaler were then fired to waste using an automated method and actuations 119 and 120 (EoU) collected.

The results are quoted in Table 3 as the mean dose for 10 inhalers as a percentage of the nominal dose of the formulation. At the end of use the data is quoted as the variability of the mean dose across the 10 inhalers.

The data shows that the percentage of the target dose is increased in the formulations of the present invention. Furthermore, the variability of the dose dispensed (at the end

of use) for the formulations of the invention which contain a surfactant is reduced as can be seen by the reduction in the percentage relative standard deviation.

SAMPLE	Beginning of Use Dose (% Target)	End of Use Dose Variability (% RSD)
CONTROL 1 Surfactant-free	78.8 %	4.0 %
EXAMPLE 1	89.6 %	2.9 %
CONTROL 2 Surfactant-free	85.2 %	7.5 %
Example 2	96.0 %	2.2 %
CONTROL 3 DRUG 1 Surfactant-free	80.7 %	5.4 %
Example 3 DRUG 1	88.1%	2.1 %
CONTROL 3 DRUG 2 Surfactant-free	95.7 %	4.4 %
Example 3 DRUG 2	97.9 %	2.4 %

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